

(12) INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(19) World Intellectual Property Organization
International Bureau



(43) International Publication Date
24 December 2003 (24.12.2003)

PCT

(10) International Publication Number
WO 03/105716 A1

(51) International Patent Classification⁷: A61C 7/08, A63B 71/08

(21) International Application Number: PCT/GB03/02566

(22) International Filing Date: 13 June 2003 (13.06.2003)

(25) Filing Language: English

(26) Publication Language: English

(30) Priority Data:
0213584.6 13 June 2002 (13.06.2002) GB

(71) Applicant and

(72) Inventor: MOHINDRA, Naresh, Kumar [GB/GB]; 18 Wimpole Street, London W1M 7AD (GB).

(74) Agents: BUCKLEY, Guy, Julian et al.; Castles, 1 King Street, Bakewell, Derbyshire DE45 1DZ (GB).

(81) Designated States (*national*): AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NI, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW.

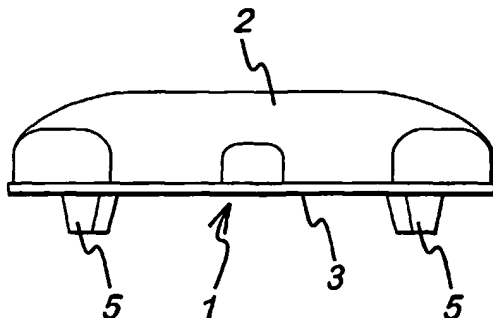
(84) Designated States (*regional*): ARIPO patent (GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, RO, SE, SI, SK, TR), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG).

Published:

— with international search report

For two-letter codes and other abbreviations, refer to the "Guidance Notes on Codes and Abbreviations" appearing at the beginning of each regular issue of the PCT Gazette.

(54) Title: REDUCING FACIAL AGEING AND APPLIANCE THEREFOR



(57) Abstract: A mass produced universal dental appliance (1) comprises two parts, the parts intended in use to contact the posterior teeth on respective opposite sides of either the upper or lower jaw, each part comprising a composite structure comprising: iii) a first layer (2) formed a durable, resilient, elastomeric material having a softening point in the range from 35 to 100°C and which in use contacts and grips the occlusal biting surfaces of the posterior teeth; and iv) a second layer (3) formed from a durable, non-deformable material having a softening point over 100°C and which in use provides a bite plate; wherein the second layer of each part is provided with a protrusion (5), formed of durable, non-deformable material having a softening point over 100°C, which extends from at least 2 mm up to 20 mm from the

surface (4) of the bite plate away from the first layer and which is positioned such that in use the protrusion extends from the surface of the bite plate above at least a part of the first and/or second molar teeth which are in contact with the first layer. The appliance is useful in a method for reducing facial aging.

WO 03/105716 A1

Reducing facial ageing and appliance therefor

This invention is concerned with a method of reducing facial aging and an appliance which is intended to be worn in the mouth during an exercise programme which, over time, can lead to a reduction in facial aging.

Pivot appliances have been used in dentistry since the 1930s to alleviate pain experienced by patients suffering from misaligned jaws, caused by inclines of the teeth. The original purpose of the pivot appliance was to separate the jaws so that inclines of the teeth would not dictate how the jaws met and thereby allow the bite of the patient to be adjusted to a more comfortable position. Use of the appliance on a temporary basis would allow the facial muscles to relax, resulting in the jaw and the condyle in the temporo mandibular joint (tmj) resting in an unrestrained position. This consequently would result in relief of pain associated with the tmj disorder.

The pivot appliance was made from a plaster mould of the patient's mouth, which mould was made by taking an impression of the lower teeth with a dental impression material. From this impression, a mould was made in plaster, which was an exact duplicate of the patient's lower jaw. This mould was then used to make a pivot appliance from a rigid moulding material, which would include wire clips to fit the appliance in the patient's mouth. The optimal thickness of the moulding material of the appliance that provided the biting surface was determined e.g. by using the command swallowing technique to establish the correct biting position for the particular patient. The patient would wear the fitted appliance under the direction and supervision of a dentist for such time until relief of pain was achieved. The appliance was then removed. The problem with this temporary procedure was that it sometimes tended to provide only temporary relief: after the appliance had been removed, there was a tendency for the jaws to return over time back to their original, painful biting position.

Some dentist would advocate that the bite of the patient had to be permanently altered to achieve permanent relief from tmj disorder. In such circumstances, orthodontic treatment was usually employed, to change the position of the teeth or by crowning the teeth.

In 1996, Dr N. K. Mohindra published a paper, in the British Dental Journal, entitled "A preliminary report on determining the vertical height of occlusion by the position of the mandible in the swallowing technique". In this paper, Dr Mohindra reported that a pivotal appliance could be used to increase the vertical dimension significantly, e.g. by up to 19 mm, beyond the normal resting position of the mandible without patients experiencing problems. Prior to this report, doctors and dentists had considered that the vertical dimension should not be increased by more than about 2-3 mm and definitely not beyond the resting position of the jaw. The appliance used in these experiments was made in a laboratory by a trained and approved dental technician.

10

In 2002, Dr Mohindra published a second paper in the British Dental Journal, entitled "The effect of increasing vertical dimension on facial aesthetics". In this paper, Dr Mohindra reported that 80% of patients whose vertical dimension had been increased permanently by the use of a pivot appliance thought they looked between 5 and 20 years younger, and that these views were backed up by an independent panel who studied before and after photographs of the patients and reached same conclusions.

15

Subsequently, Dr Mohindra developed a facial rejuvenator which improved facial aesthetics without permanently increasing vertical dimension of occlusion. The rejuvenator, like the earlier dental appliance, required to be custom made for each patient in a laboratory by a trained dental technician. The rejuvenator comprised a substantially U-shaped layer formed from a durable, non-deformable material having a softening point over 100°C which was custom moulded to fit over all the teeth on the lower jaw and which in use provided a bite plate. Two projections extended from the surface of the bite plate and were positioned on the bite plate over at least a part of the first molar tooth on both sides of the jaw. The projections were made of a durable, non-deformable material having a softening point over 50°C, and were custom moulded to the vertical height of occlusion for each patient, as determined by the command swallowing technique.

20

25

The rejuvenator was removable and so only increases the vertical dimension of occlusion for the short period of time when the appliance is in the mouth of the patient. The rejuvenator was based on the original pivot appliance and is made in the same way, i.e. in a laboratory by a trained and approved dental technician.

30

US-6415794 and US-A-6539943 disclose a dental appliance for use by athletes during periods of exertion. The appliance consists of an occlusal posterior pad made of quadruple composite material comprising four layers of distinct materials, further comprising a first layer of a durable, resilient material, a second layer of non-softenable, flexible material, a third layer of a hard, very durable material, and a fourth layer of softenable material, engageable with the occlusal surfaces to space apart the upper and lower teeth, to absorb shock and clenching stress. An adjustable arch adapted to expand and contract to be moulded to the palate is provided connecting the posterior pads together with the mouth and out of the way of the tongue to maintain the position of the occlusal posterior pads within the mouth during use and to prevent loss of the pads such as by swallowing. The appliance may be fitted using a boil and bite technique, for example by a doctor or dentist, with no requirement for customized laboratory moulding processes.

US-A-6092523 discloses an anti-snoring device having a dental overlay portion and a guide ramp portion slidably mounted in the dental portion. The device may be fitted using a boil and bite technique, for example by a doctor or dentist, with no requirement for customized laboratory moulding processes.

The object of the present invention is to provide an appliance for reducing facial aging that can be mass produced on an industrial basis, thereby obviating the hitherto necessity for customized manufacture in a laboratory of an appliance that has been individually designed and made for a particular patient.

The present invention is as set out in the accompanying claims.

In accordance with one aspect of the present invention there is provided a mass produced universal dental appliance suitable for use in a method of reducing facial aging, which appliance comprises two parts, the parts intended in use to contact the posterior teeth on respective opposite sides of either the upper or lower jaw, each part comprising a composite structure comprising:

- i) a first layer formed a durable, resilient, elastomeric material having a softening point in the range from 35 to 100°C and which in use contacts and grips the occlusal biting surfaces of the posterior teeth; and

- ii) a second layer formed from a durable, non-deformable material having a softening point over 100°C and which in use provides a bite plate;
wherein the second layer of each part is provided with a protrusion, formed of durable, non-deformable material having a softening point over 100°C, which extends from at least 2mm up to 20mm from the surface of the bite plate away from the first layer and which is positioned such that in use the protrusion extends from the surface of the bite plate above at least a part of the first and/or second molar teeth which are in contact with the first layer.
- 10 The universal appliance is adapted to provide a predetermined vertical separation of the jaws determined by the command swallowing technique and, advantageously, can be either fitted by e.g. a doctor or dentist without requiring use of customized laboratory processes or a dental technician, or it may be purchased over-the-counter and fitted by the individual user.
- 15 The universal appliance is intended to be used by a patient to reduce the signs of facial aging. The universal appliance advantageously does not have to be made individually for a patient, unlike the rejuvenator.
The appliance may be worn at any time of the day, when the patient is awake or asleep. Preferably, the appliance is worn during eating or during sleeping.
- 20 Use of the universal appliance will be generally prescribed by a doctor or a dentist or, in the case of an over-the-counter purchase, as prescribed on the accompanying instructions for use.
- The appliance is preferably worn for from about 3 to about 12 hours in any day. It is recommended not to wear the appliance for 24 hours of the day.
- 25 The appliance is preferably used over a continuous period of from 4 to 10 weeks, typically 6 weeks, with a preferred interval before reuse of from 3 to 6 months e.g. 4 months.
- 30 The appliance of the present invention is shaped to fit over at least the biting surface of the posterior teeth of the upper or lower jaw, preferably the lower jaw. It is preferred that the appliance is substantially U-shaped, so as to fit comfortably over both the anterior and posterior teeth. Alternatively, the appliance may comprise two separate portions which fit

over only the posterior teeth on either side of the respective jaw, with a bridging means to connect the two portions.

Each part of the two parts of the appliance of the present invention which fit over the posterior teeth on both sides of the upper or lower jaw consists of a composite structure comprising

- i) a first layer formed a durable, resilient, elastomeric material having a softening point in the range from about 35 to 100°C and which in use contacts and grips the occlusal biting surfaces of the posterior teeth; and
- ii) a second layer formed from a durable, non-deformable material having a softening point over 100°C and which in use provides a bite plate.

The second layer is provided with a protrusion formed of durable, non-deformable material having a softening point over 100°C which extends from at least about 2mm up to about 20mm, preferably from about 5 mm up to about 15 mm, most preferably from about 7mm to about 10mm e.g. 9 mm, from the surface of the bite plate away from the first layer and which is positioned such that in use it is on the bite plate above at least a part of the first and/or second molar teeth which are in contact with the first layer.

The protrusion is preferably formed integrally with the second layer.

Preferably, the protrusion is centrally located above at least a part of the first and/or second molar teeth. More preferably, the width of the protrusion is less than the width of the molar(s) above which the protrusion is intended to be positioned.

The shape of the protrusion is not important, provided that in use it is comfortable for the patient and when the appliance is fitted the protrusion provides a point above the first and/or second molars on the lower jaw about which the lower jaw may pivot, if forced to do so.

In another aspect, the present invention provides a method of reducing facial aging, which method comprises fitting an appliance as described above on either the upper or lower jaw, preferably the lower jaw, and exercising the lower jaw by repeatedly dropping the lower jaw and then lifting the lower jaw and closing it against the upper jaw with the appliance between the teeth.

Exercising may be undertaken when the patient is conscious or asleep. Exercising may take place actively, for example during eating or at a time when the patient deliberately exercises. Alternatively, exercising may take place passively, for example when the patient is asleep or simply performing normal daytime activities. In such passive exercise, the facial muscles are stretched by the jaw adopting a new resting position.

Exercising is preferably achieved by wearing the appliance continuously for from about 3 to about 12 hours in any day (it is recommended not to wear the appliance for 24 hours of the day). Exercising is preferably undertaken on a daily basis over a period of from 4 to 10 weeks, typically 6 weeks. A break from exercising of from 3 to 6 months e.g. 4 months is preferably taken before commencing another period of exercising.

The first layer of the appliance is formed from a durable, resilient, elastomeric material, preferably having a softening point in the range from about 35 to 100°C, preferably about 35 to 50°C, more preferably from about 35 to 40°C. Such materials are well known in the art and are commonly used in the manufacture of boil and bite type dental products, such as those described in US-A-6092253, US-A-6415794 and US-A-6539943. Examples of suitable materials are styrene block copolymers, polyolefin rubbers, acrylate based elastomers and ethylene vinyl acetate copolymers, and mixtures thereof. Commercially available materials include Elvax and Engage available from DuPont, Kraton thermoplastic rubber available from Shell, Santoprene available from Advanced Elastomer Systems and Dynaflex available from GLS. The material used to form the first layer must become mouldable when placed in water at or close to boiling temperatures.

The first layer is preferably from 5 to 15 mm thick, more preferably 8 to 12 mm thick, before fitting. After fitting, the thickness of the layer will vary from point to point along the length appliance. Preferably, after fitting, the thickness of the first layer does not go below 1 mm.

It is the employment of the first layer material in the composite structure of the appliance that enables the appliance during fitting to adjust to provide a vertical separation of the jaws determined by the command swallow technique.

The second layer of the appliance is formed from a durable, non-deformable material having a softening point over 100°C, e.g. 150°C or more. Such materials are well known in the art and are commonly used in the manufacture of boil and bite type dental products, such as those described in US-A-6092253, US-A-6415794 and US-A-6539943. Examples of suitable materials include polycarbonate resins, high density polyethylene and polypropylene and methylmethacrylate based thermoplastics. Commercially available materials include Escorene HD-6706 available from Exxon and AP6112-HS available from Huntsman. The material used to make the second layer must not become softened in boiling water.

- 10 The second layer is preferably from about 1 to about 15 mm thick, preferably from about 3 to about 9 mm thick. The thickness of the second layer is not affected by fitting.

The second layer is provided with a protrusion formed of durable, non-deformable material having a softening point over 100°C which extends from at least about 2mm up to about 20mm, preferably from about 5 mm up to about 15 mm, most preferably from about 7mm to about 10mm e.g. 9 mm, from the surface of the bite plate away from the first layer and which is positioned such that in use it is on the bite plate above at least a part of the first and/or second molar teeth which are in contact with the first layer.

- 20 The appliance may comprise a third layer of material located between the first and second layers. If present, such a third layer is preferably formed of a durable resilient material having a softening point above 100°C, preferably above 150°C. Such suitable materials are mentioned above.

- 25 The protrusion is preferably formed out of the same material as the second layer and is preferably formed integrally with the second layer. Together, the protrusion and second layer are preferably no more than 22 mm thick at their thickest point, more preferably no more than 15mm thick at their thickest point.

- 30 The appliance of the present invention can be fitted to a patient employing a similar boil and bite procedure as disclosed in US-A-6415794, US-A-6539943 or US-A-6092523. The fitting should include the use of the command swallow technique.

The appliance of the present invention is useful for reducing facial aging. The aging of the face basically involves two factors. These are intrinsic and extrinsic factors. The intrinsic factors basically involve atrophy i.e. the reduction in number of cells for instance by the age of 60 (typically, we only have 60% of the muscle cells that we had when we were in our 20's).

5 The extrinsic factors involve damage done to cells by environmental factors e.g. sun, smoke, toxins produced by bacteria and viruses. The process involved is basically a form of chronic inflammation. Both these processes (cell death and chronic inflammation) in their early stages are reversible. Exercising with the appliance of the present invention can help reverse these processes in their early stages. Accordingly, all diseases, which are related to facial aging or
10 inflammatory conditions, could be alleviated to some extent by exercising with the appliance. For example, exercising with the appliance could help to alleviate, to some extent, some of the symptoms associated with suffers of Alzheimer's, chronic sinusitis, age related deterioration in eyesight, tangelacetasis, solar damage to the skin, acne, and bacterial infections, such as ear infections. This list is not exhaustive.

15

The invention in its various embodiments shall now be further described by way of exemplification with reference to the accompanying drawings, in which:

Fig 1 is a plan view of an appliance in accordance with the present invention.

20 Fig 2 is a cross-sectional view of the appliance shown in Fig 1 along the line A-A.

Fig 3 is an end view of the appliance shown in Fig 1 along the line B-B.

Fig 4 is an end view of the appliance shown in Fig 1 along the line C-C.

Fig5 is a view from above of another appliance in accordance with the present invention.

Fig 6 is an end view of the appliance shown in Fig 5 along the line D-D.

25 Fig 7 is a view from below of the appliance shown in Fig 5.

Fig 8 is a cross-sectional view of the appliance shown in Fig 5 along the line E-E.

A U-shaped universal facial rejuvenator appliance 1, as shown in Figs 1-8, may comprise a first layer 2, about 10 mm thick formed of a commercially available substantially transparent
30 elastomeric EVA copolymer having a softening point of about 36°C, and a second layer 3, about 1.5 mm thick formed of a commercially available substantially transparent polycarbonate having a softening point of about 190°C. The surface 4 of the second layer 3 remote from the first layer 2 forms, in use, a bite plate. Extending from the surface of the bite

plate 4 are two projections 5 which may be formed integrally with the second layer. The projections extend approximately 3 mm above the surface 4.

5 The first layer 2 and second layer 3 may be adhered together, with the use of an appropriate adhesive, or may be heat formed together. If heat forming is employed, the second layer may be provided with a plurality small orifices 6 or projections into which or around which the first layer engages to secure itself in position.

10 The universal appliance 1 can be fitted to a patient by initially heating the appliance by submerging it in near boiling water for about 30 seconds, or such other time so as to render the material of the first layer 2 mouldable. The patient is required to open the mouth and the appliance is then placed over the teeth on the patient's lower jaw, with the first layer 2 in contact with the teeth and the two projections 5 positioned over both first molar teeth. The patient is then required to gently raise the lower jaw until the teeth on the upper jaw contact
15 the top of the projections 5. The patient then closes their lips and swallows. The pressure applied to the to the appliance 1 by swallowing causes the material of the first layer 2 to deform and mould itself to the shape of the teeth on the lower jaw. The appliance can then be carefully removed from the mouth of the patient and submerged in cold water to accelerate the cooling of the appliance to ambient. Once the appliance has cooled, any excess of the first
20 layer material can be trimmed away until it forms a comfortable fit.

The fitting of the appliance can readily be performed by a doctor or dentist, without the services of a dental technician or having to resort to custom moulding practices in a laboratory, or by the patient without third party assistance.

25

Once the appliance 1 has cooled to ambient temperatures and been trimmed, it may be used by the patient for reducing facial aging.

30 The process for reducing facial aging can commence as soon as the appliance has cooled to a temperature where the material used to form the first layer has solidified sufficiently for use of the appliance not to cause remoulding of that layer. The patient positions the appliance 1 in the mouth, preferably just before eating a meal. The patient then exercises the lower jaw, e.g. during eating or sleeping, by raising the lower jaw thereby to bring the teeth on the upper jaw

into contact with the protrusions and then closing the teeth on the upper and lower jaw around the appliance. In so doing, the lower jaw is forced to pivot slightly about the protections and cause the facial muscles controlling the lower jaw to work harder. The muscles are then permitted to relax and the lower jaw is dropped. The exercise is preferably repeated for about
5 6 to 7 hours a day over a period of about 6 weeks.

The universal appliance may be tested on a sample of patients. The patients may be asked to wear the appliance at mealtimes over a specified period of time. 12 patients wore the appliance for 6 weeks, 1 for 7 weeks, 4 for 8 weeks, 2 for 9 weeks, 3 for 10 weeks, and 6 for
10 12 weeks. The results are as follows:

The patients were asked:

- 1) How comfortable was the appliance in the mouth on a scale of 0 to 3 (0 being very uncomfortable and 3 being very comfortable)?
15 4 patients thought it was 1 on the scale of 0 to 3, 15 thought 2, and 9 thought 3.
- 2) If you suffered from neckaches, headaches, shoulder pains, did these improve?
4 patients said they suffered from neck ache and 100% reported an improvement. 5 suffered from shoulder pain and 80% reported an improvement. 3 had headaches and 66% reported improvement.
- 20 3) Do you think the treatment enhanced your facial features? Give your response on a scale between 0 and 3, 0 being no improvement and 3 being good. The number of features listed was 11 and were as follows: skin above the upper eyelid, size of the eyes, crows feet, lateral droop of the eyes, bags under the eyes, improvement in skin, naso labial folds, lips, jaw line, cheeks. The results were as follows;
25 On an average patients noted an improvement in 7.5 of the 11 features:
skin above the upper eyelid: 60.7% noted an improvement; size of the eyes: 57.1%; crows feet: 64.3%; lateral droop of the eyes: 71% noted an improvement; bags under the eyes: 57.1%; skin: 85.7%; naso labial folds: 71.4%; lips:89.3%; jaw line:64.3%; neck: 64.3%; cheeks:71.4%.
- 30 All these improvements were from mild to very good, suggesting that the universal appliance could be successfully employed to reduce facial aging.

Claims:

1. A mass produced universal dental appliance (1) suitable for use in a method of reducing facial aging, which appliance comprises two parts, the parts intended in use to contact the posterior teeth on respective opposite sides of either the upper or lower jaw, each part comprising a composite structure comprising:
- 5
- i) a first layer (2) formed a durable, resilient, elastomeric material having a softening point in the range from 35 to 100°C and which in use contacts and grips the occlusal biting surfaces of the posterior teeth; and
 - 10 ii) a second layer (3) formed from a durable, non-deformable material having a softening point over 100°C and which in use provides a bite plate;
- wherein the second layer of each part is provided with a protrusion (5), formed of durable, non-deformable material having a softening point over 100°C, which extends from at least 2mm up to 20mm from the surface (4) of the bite plate away from the first layer and
- 15 which is positioned such that in use the protrusion extends from the surface of the bite plate above at least a part of the first and/or second molar teeth which are in contact with the first layer.
2. An appliance as claimed in claim 1, wherein the protrusion is formed integrally with the
- 20 second layer.
3. An appliance as claimed in claim 1 or claim 2, wherein the protrusion is centrally located above at least a part of the first and/or second molar teeth.
- 25 4. An appliance as claimed in any one of the preceding claims, wherein the width of the protrusion is less than the width of the molar teeth above which the protrusion is intended to be positioned.
- 30 5. A method for reducing facial aging in an individual, which method comprises fitting an appliance as claimed in any one of the preceding claims over at least the posterior teeth on both sides of either the upper jaw or lower jaw of an individual, and exercising the lower jaw by repeatedly dropping the lower jaw and then lifting the lower jaw and forcing it to close against the upper jaw with the appliance between the teeth.

6. A method as claimed in claim 6, wherein said exercising takes place for at least 3 hours per day, over a period of at least 4 weeks.

7. A method as claimed in claim 5 or claim 6, wherein said exercise takes place actively.

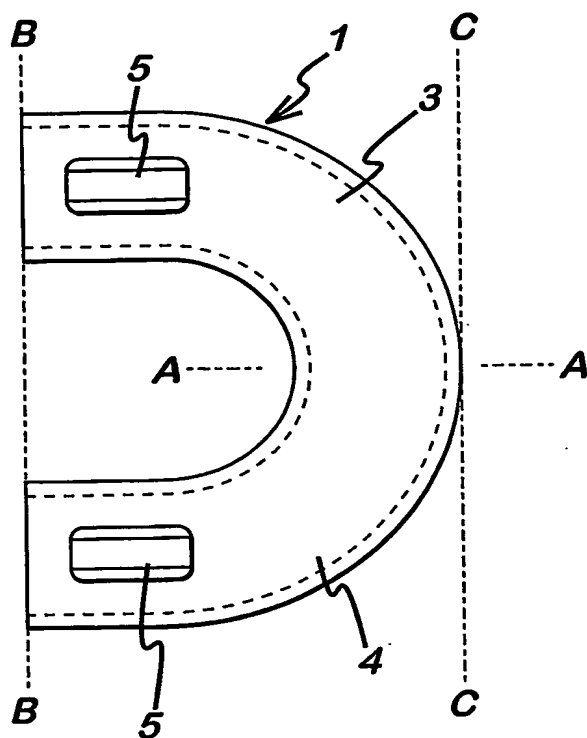
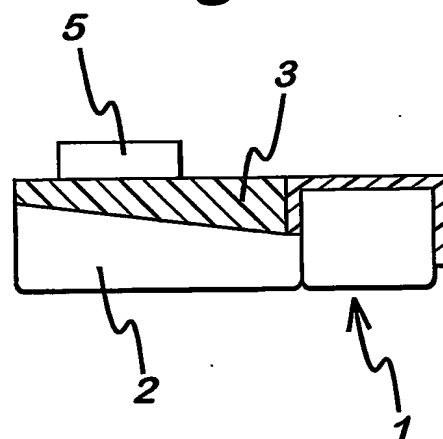
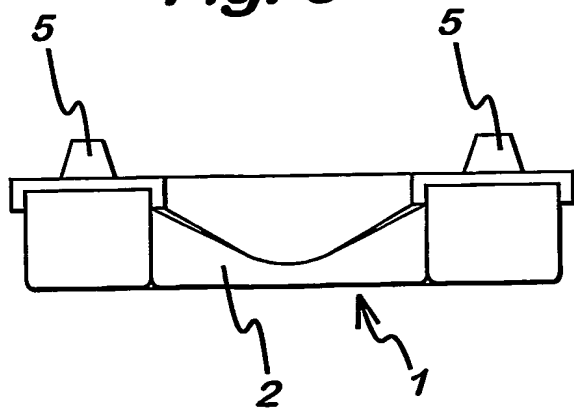
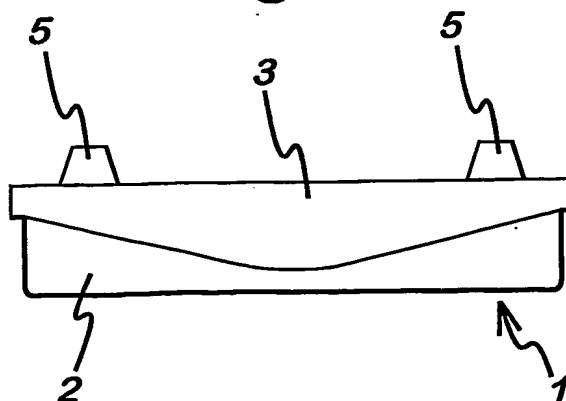
5

8. A method as claimed in claim 7, wherein said exercising takes place when eating.

9. A method as claimed in claim 5 or claim 6, wherein said exercising takes place passively.

10

1/2

Fig. 1**Fig. 2****Fig. 3****Fig. 4**

2/2

Fig. 5

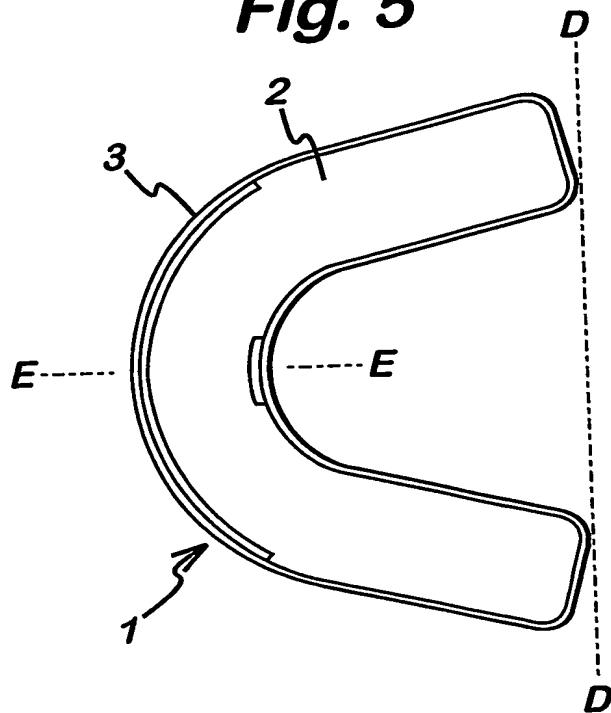


Fig. 6

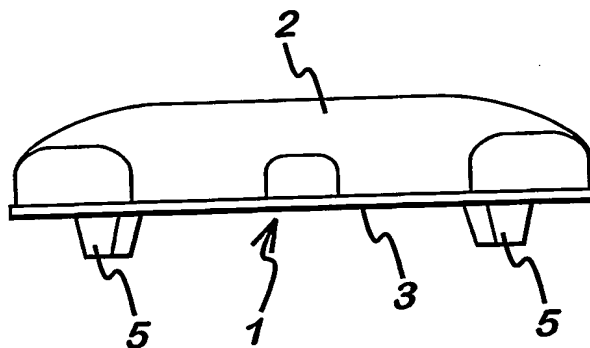


Fig. 7

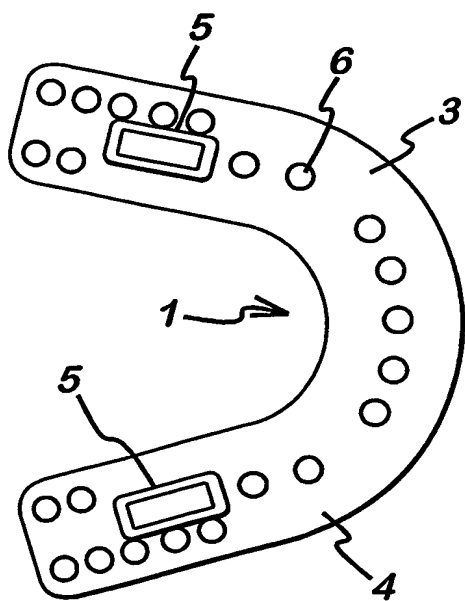
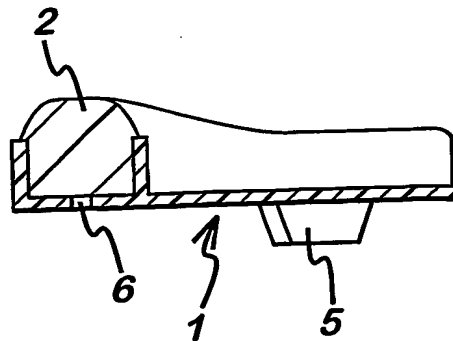


Fig. 8



INTERNATIONAL SEARCH REPORT

Internat Application No
PCT/GB 03/02566

A. CLASSIFICATION OF SUBJECT MATTER
IPC 7 A61C7/08 A63B71/08

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)
IPC 7 A61C A63B A61F

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the International search (name of data base and, where practical, search terms used)

EPO-Internal, WPI Data, PAJ

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	US 5 299 936 A (UENO) 5 April 1994 (1994-04-05) column 2, line 16 - line 53 column 6, line 59 -column 7, line 14 claim 4; figures 7-13	1-4
A	WO 00 35369 A (FARRELL) 22 June 2000 (2000-06-22) the whole document	1
A	US 5 879 155 A (KITTELSEN) 9 March 1999 (1999-03-09) column 6, line 62 -column 7, line 35 figures 17,18	1
A	US 6 371 758 B1 (KITTELSEN) 16 April 2002 (2002-04-16) the whole document	1

☐ Further documents are listed in the continuation of box C.

☒ Patent family members are listed in annex.

* Special categories of cited documents :

- *A* document defining the general state of the art which is not considered to be of particular relevance
- *E* earlier document but published on or after the international filing date
- *L* document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)
- *O* document referring to an oral disclosure, use, exhibition or other means
- *P* document published prior to the international filing date but later than the priority date claimed

- *T* later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention
- *X* document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone
- *Y* document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.
- *G* document member of the same patent family

Date of the actual completion of the international search

20 August 2003

Date of mailing of the international search report

28/08/2003

Name and mailing address of the ISA

European Patent Office, P.B. 5818 Patentlaan 2
NL - 2280 HV Rijswijk
Tel. (+31-70) 340-2040, Tx. 31 651 epo nl,
Fax: (+31-70) 340-3016

Authorized officer

Raybould, B

INTERNATIONAL SEARCH REPORT

Inter application No.
PCT/GB 03/02566

Box I Observations where certain claims were found unsearchable (Continuation of item 1 of first sheet)

This International Search Report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:

1. ☒ Claims Nos.: 5-9
because they relate to subject matter not required to be searched by this Authority, namely:
Rule 39.1(iv) PCT - Method for treatment of the human or animal body by therapy
2. ☐ Claims Nos.:
because they relate to parts of the International Application that do not comply with the prescribed requirements to such an extent that no meaningful International Search can be carried out, specifically:
3. ☐ Claims Nos.:
because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a).

Box II Observations where unity of invention is lacking (Continuation of item 2 of first sheet)

This International Searching Authority found multiple inventions in this International application, as follows:

1. ☐ As all required additional search fees were timely paid by the applicant, this International Search Report covers all searchable claims.
2. ☐ As all searchable claims could be searched without effort justifying an additional fee, this Authority did not invite payment of any additional fee.
3. ☐ As only some of the required additional search fees were timely paid by the applicant, this International Search Report covers only those claims for which fees were paid, specifically claims Nos.:
4. ☐ No required additional search fees were timely paid by the applicant. Consequently, this International Search Report is restricted to the invention first mentioned in the claims; it is covered by claims Nos.:

Remark on Protest

- ☐ The additional search fees were accompanied by the applicant's protest.
☐ No protest accompanied the payment of additional search fees.

INTERNATIONAL SEARCH REPORT

Internat
pplication No
PCT/GB 03/02566

Patent document cited in search report		Publication date	Patent family member(s)	Publication date
US 5299936	A	05-04-1994	JP 5058174 U	03-08-1993
			JP 7039809 U	18-07-1995
			JP 6015610 U	01-03-1994
			FR 2681523 A1	26-03-1993
			GB 2259649 A , B	24-03-1993
			US 5513984 A	07-05-1996
WO 0035369	A	22-06-2000	AU 6452799 A	03-07-2000
			WO 0035369 A1	22-06-2000
			CA 2393681 A1	22-06-2000
			EP 1139905 A1	10-10-2001
			US 2003019497 A1	30-01-2003
US 5879155	A	09-03-1999	US 5836761 A	17-11-1998
			US 6200133 B1	13-03-2001
			US 6371758 B1	16-04-2002
US 6371758	B1	16-04-2002	US 5836761 A	17-11-1998
			US 6200133 B1	13-03-2001
			US 5879155 A	09-03-1999